

Welcome to STN International! Enter x:x

LOGINID: ssspta1653hxp

PASSWORD:

TERMINAL (ENTER 1, 2, 3, OR ?):2

\* \* \* \* \* \* \* \* \* \* \* \* \* \* \* Welcome to STN International \* \* \* \* \* \* \* \* \* \* \*

NEWS 1 Web Page URLs for STN Seminar Schedule - N. America  
NEWS 2 Apr 08 "Ask CAS" for self-help around the clock  
NEWS 3 Apr 09 BEILSTEIN: Reload and Implementation of a New Subject Area  
NEWS 4 Apr 09 ZDB will be removed from STN  
NEWS 5 Apr 19 US Patent Applications available in IFICDB, IFIPAT, and

IFIUDB NEWS 6 Apr 22 Records from IP.com available in CAPLUS, HCAPLUS, and

ZCAPPLUS  
NEWS 7 Apr 22 BIOSIS Gene Names now available in TOXCENTER  
NEWS 8 Apr 22 Federal Research in Progress (FEDRIP) now available  
NEWS 9 Jun 03 New e-mail delivery for search results now available  
NEWS 10 Jun 10 MEDLINE Reload  
NEWS 11 Jun 10 PCTFULL has been reloaded  
NEWS 12 Jul 02 FOREGE no longer contains STANDARDS file segment  
NEWS 13 Jul 19 NTIS to be reloaded July 28, 2002

NEWS 14 Jul 22 USAN to be reloaded July 28, 2002;  
saved answer sets no longer valid  
NEWS 15 Jul 29 Enhanced polymer searching in REGISTRY  
NEWS 16 Jul 30 NETFIRST to be removed from STN

NEWS EXPRESS	February 1 CURRENT WINDOWS VERSION IS V6.0d, CURRENT MACINTOSH VERSION IS V6.0a(ENG) AND V6.0Ja(JP), AND CURRENT DISCOVER FILE IS DATED 05 FEBRUARY 2002
NEWS HOURS	STN Operating Hours Plus Help Desk Availability
NEWS INTER	General Internet Information
NEWS LOGIN	Welcome Banner and News Items
NEWS PHONE	Direct Dial and Telecommunication Network Access to STN
NEWS WWW	CAS World Wide Web Site (general information)

Enter NEWS followed by the item number or name to see news on that specific topic.

All use of STN is subject to the provisions of the STN Customer agreement. Please note that this agreement limits use to scientific research. Use for software development or design or implementation of commercial gateways or other similar uses is prohibited and may result in loss of user privileges and other penalties.

=> file medline, biosis

COST IN U.S. DOLLARS

SINCE FILE

TOTAL

FULL ESTIMATED COST

ENTRY

SESSION

0.21

0.21

FILE 'MEDLINE' ENTERED AT 17:43:02 ON 07 AUG 2002

FILE 'BIOSIS' ENTERED AT 17:43:02 ON 07 AUG 2002  
COPYRIGHT (C) 2002 BIOLOGICAL ABSTRACTS INC. (R)

=> s angiogenesis

L1 29859 ANGIOGENESIS

=> s 12 and inhibition

L2 NOT FOUND

The L-number entered could not be found. To see the definition  
of L-numbers, enter DISPLAY HISTORY at an arrow prompt (>).

=> s 11 and inhibition

L2 4274 L1 AND INHIBITION

=> s 12 and method

L3 540 L2 AND METHOD

=> s 13 and composition

L4 4 L3 AND COMPOSITION

=> d 14 ti abs ibib tot

L4 ANSWER 1 OF 4 BIOSIS COPYRIGHT 2002 BIOLOGICAL ABSTRACTS INC.

TI Pharmaceutical compositions and methods of inhibiting **angiogenesis**  
using naaladase inhibitors.

AB The present disclosure relates to a **method** of inhibiting  
**angiogenesis** comprising administering a N-Acetylated alpha-Linked  
Acidic Dipeptidase (NAALADase) inhibitor to a patient in need thereof,  
and  
a pharmaceutical **composition** comprising an anti-angiogenic  
effective amount of a NAALADase inhibitor and a pharmaceutically  
acceptable carrier.

ACCESSION NUMBER: 2002:358717 BIOSIS

DOCUMENT NUMBER: PREV200200358717

TITLE: Pharmaceutical compositions and methods of inhibiting  
**angiogenesis** using naaladase inhibitors.

AUTHOR(S): Slusher, Barbara S.; Lapidus, Rena G. (1)

CORPORATE SOURCE: (1) Pikesville, MD USA  
ASSIGNEE: Guilford Pharmaceuticals Inc., Baltimore, MD,  
USA

PATENT INFORMATION: US 6395718 May 28, 2002

SOURCE: Official Gazette of the United States Patent and Trademark  
Office Patents, (May 28, 2002) Vol. 1258, No. 4, pp. No  
Pagination. <http://www.uspto.gov/web/menu/patdata.html>.  
e-file.

ISSN: 0098-1133.

DOCUMENT TYPE: Patent  
LANGUAGE: English

L4 ANSWER 2 OF 4 BIOSIS COPYRIGHT 2002 BIOLOGICAL ABSTRACTS INC.  
TI Methods and compositions for inhibiting angiogenesis.  
AB The present invention provides a **method** of inhibiting angiogenesis within a tissue by providing exogenous SLED to cells associated with the tissue. The presence of exogenous SLED inhibits angiogenesis within the tissue, in part by interfering with the ability of vascular endothelia to expand within the tissue. The invention also provides a **method** for determining the severity of a tumor by assaying for the presence of SLED within the tumor. To facilitate the inventive methods, the present invention provides pharmaceutical compositions including sources of SLED.

ACCESSION NUMBER: 2001:521933 BIOSIS

DOCUMENT NUMBER: PREV200100521933

TITLE: Methods and compositions for inhibiting angiogenesis.

AUTHOR(S): Bouck, Noel P. (1); Dawson, David W.; Gillis, Paul R.

CORPORATE SOURCE: (1) Oak Park, IL USA

ASSIGNEE: Northwestern University

PATENT INFORMATION: US 6288024 September 11, 2001

SOURCE: Official Gazette of the United States Patent and Trademark Office Patents, (Sep. 11, 2001) Vol. 1250, No. 2, pp. No Pagination. e-file.

ISSN: 0098-1133.

DOCUMENT TYPE: Patent

LANGUAGE: English

L4 ANSWER 3 OF 4 BIOSIS COPYRIGHT 2002 BIOLOGICAL ABSTRACTS INC.

TI Inhibition of hair growth.

AB A **method** of inhibiting hair growth in a mammal includes applying, to an area of skin from which reduced hair growth is desired, a dermatologically acceptable **composition** containing a non-steroidal suppressor of angiogenesis.

ACCESSION NUMBER: 2001:188368 BIOSIS

DOCUMENT NUMBER: PREV200100188368

TITLE: Inhibition of hair growth.

AUTHOR(S): Ahluwalia, Gurpreet S. (1); Styczynski, Peter; Shander, Douglas

CORPORATE SOURCE: (1) 8632 Stable View Ct., Gaithersburg, MD, 20879 USA

PATENT INFORMATION: US 6093748 July 25, 2000

SOURCE: Official Gazette of the United States Patent and Trademark Office Patents, (July 25, 2000) Vol. 1236, No. 4, pp. No Pagination. e-file.

ISSN: 0098-1133.

DOCUMENT TYPE: Patent

LANGUAGE: English

L4 ANSWER 4 OF 4 BIOSIS COPYRIGHT 2002 BIOLOGICAL ABSTRACTS INC.

TI Alteration of ganglioside **composition** by stable transfection with antisense vectors against GD3-synthase gene expression.

AB Gangliosides are ubiquitous components of mammalian cells. Their expression is frequently altered in many tumor types. We previously showed

that alteration of the ganglioside **composition** often resulted in changes in cellular morphology and differentiation of cultured cells. In this study, we targeted sialyltransferase gene expression by the antisense

knockdown experiment, and the results showed that inhibition of the expression of gangliosides GD3 and O-acetylated GD3 (OAc-GD3) in the neuroblastoma F-11 cells greatly reduced the tumor growth in nude mice.

The sense and antisense vectors containing either a 5' end fragment or the entire sequence of the cDNA coding for GD3-synthase were prepared and used

in separate experiments to transfet the F-11 cells which express high

levels of gangliosides GD3 and OAc-GD3. Single clones were isolated and expanded. Both the activity of the GD3-synthase and the concentrations of GD3 and OAc-GD3 in the antisense-transfected cells were dramatically decreased as a result of transfection with the antisense expression vectors. Further characterization of the antisense-transfected cells showed reduced rates of cell growth and neurite formation and changes in cellular morphology. When the cells were inoculated in athymic nude mice, the tumor growth rate was remarkably suppressed although the tumor incidence was not affected by the altered ganglioside **composition**. These results indicate that the tumor-associated ganglioside(s) is(are) involved in regulation of tumor growth, probably through the stimulation of **angiogenesis** of the tumor.

ACCESSION NUMBER: 1999:356557 BIOSIS

DOCUMENT NUMBER: PREV199900356557

TITLE: Alteration of ganglioside **composition** by stable transfection with antisense vectors against GD3-synthase gene expression.

AUTHOR(S): Zeng, Guichao; Li, Donna D.; Gao, Luoyi; Birkle, Stephane; Bieberich, Erhard; Tokuda, Akira; Yu, Robert K. (1)

CORPORATE SOURCE: (1) Department of Biochemistry and Molecular Biophysics, Medical College of Virginia, Virginia Commonwealth University, Richmond, VA, 23298-0614 USA

SOURCE: Biochemistry, (July 6, 1999) Vol. 38, No. 27, pp. 8762-8769.

ISSN: 0006-2960.

DOCUMENT TYPE: Article

LANGUAGE: English

SUMMARY LANGUAGE: English